WEST
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Main Menu   Search Form   Posting Counts   Show S Numbers   Edit S Numbers   Preferences   Cases
Search Results -  Terms Documents  mlb13myc 0
US Patents Full-Text Database US Pre-Grant Publication Full-Text Database JPO Abstracts Database EPO Abstracts Database Derwent World Patents Index IBM Technical Disclosure Bulletins
Search:  Refine Search
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Search History
DATE: Monday, March 11, 2002 Printable Copy Create Case
Set Name Query Hit Count Set Name result set
$DB=USPT$ ; $PLUR=YES$ ; $OP=ADJ$ $\underline{L1} \qquad \text{mlb13myc} \qquad \qquad 0 \qquad \underline{L1}$

**END OF SEARCH HISTORY** 

## (FILE 'HOME' ENTERED AT 10:57:32 ON 11 MAR 2002)

INDEX 'ADISALERTS, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DDFB, DDFU, DGENE, DRUGB, DRUGLAUNCH, DRUGMONOG2, ...' ENTERED AT 10:57:41 ON

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11 MAR 2002
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FILE WPINDEX

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INDEX 'ADISALERTS, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHOS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DDFB, DDFU, DGENE, DRUGB, DRUGLAUNCH, DRUGMONOG2, ...' ENTERED AT 10:57:41 ON

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(FILE 'HOME' ENTERED AT 09:56:56 ON 11 MAR 2002)

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## SEA MLB13MYC

- FILE BIOSIS 6
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- FILE CAPLUS 5
- FILE EMBASE 2
- FILE ESBIOBASE 1
- FILE LIFESCI 1
- FILE MEDLINE
- FILE SCISEARCH
- FILE TOXCENTER
- FILE TOXLIT

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QUE MLB13MYC

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- 29 S MLB13MYC L2
  - 8 DUP REM L2 (21 DUPLICATES REMOVED)
- 0 S L2 AND CLONE 14 L4

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L3

ANSWER 24 OF 24 DGENE COPYRIGHT 2002 DERWENT INFORMATION LTD L6 AN AAV84394 DNA **DGENE** New human frizzled-related protein and associated nucleic acid, probes, ΤI vectors - transformants, antibodies and transgenic animals, used to inhibit signalling by Wnt-family cytokines, potentially useful as tumour suppressor Aaronson S; Finch P; He X; Rubin J S IN (USSH) US DEPT HEALTH & HUMAN SERVICES. PA A1 19981203 PΙ WO 9854325 WO 1998-US10974 19980529 ΑI US 1997-50495 19970623 PRAI US 1997-50417 19970529 DTPatent English LA 1999-059840 [05] os

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L3 ANSWER 8 OF 8 CAPLUS COPYRIGHT 2002 ACS DUPLICATE 5

AN 1993:618185 CAPLUS

DN 119:218185

- TI Development of immortalized cells derived from 13DPC mouse limb buds as a system to study the effects of recombinant human bone morphogenetic protein-2 (rhBMP-2) on limb bud cell differentiation
- AU Rosen, Vicki; Capparella, Joanna; McQuaid, David; Cox, Karen; Thies, R. Scott: Song, Jeffrey: Wozney, John

Scott; Song, Jeffrey; Wozney, John CS Genet. Inst. Inc., Cambridge, MA, 02140, USA

SO Prog. Clin. Biol. Res. (1993), 383A(Limb Development and Regeneration, Pt. A), 305-15
CODEN: PCBRD2; ISSN: 0361-7742

DT Journal

LA English

- Immortalization of cells derived from 13dpc mouse embryo limb buds has resulted in the establishment of a cell population, MLB13MYC that is capable of expressing differentiated phenotype in vitro when exposed to recombinant human bone morphogenetic protein-2 (rhBMP-2). Treatment of these cells for 24-48 h with rhBMP-2 resulted in the stimulation of both alk. phosphatase activity and the ability of the cells to respond to parathyroid hormone by producing cAMP. While these parameters are not unique markers of the cartilage and bone cell phenotype, they provide some evidence that MLB13MYC cells are undergoing differentiation in vitro. The presence of bone- and cartilage-specific phenotype markers after rhBMP-2 treatment of MLB13MYC cells further supports the hypothesis that MLB13MYC are a useful model system with which to study the effects of rhBMP-2 and other growth and differentiation.
- Immortalization of cells derived from 13dpc mouse embryo limb buds has resulted in the establishment of a cell population, MLB13MYC that is capable of expressing differentiated phenotype in vitro when exposed to recombinant human bone morphogenetic protein-2 (rhBMP-2). Treatment of these cells for 24-48 h with rhBMP-2 resulted in the stimulation of both alk. phosphatase activity and the ability of the cells to respond to parathyroid hormone by producing cAMP. While these parameters are not unique markers of the cartilage and bone cell phenotype, they provide some evidence that MLB13MYC cells are undergoing differentiation in vitro. The presence of bone- and cartilage-specific phenotype markers after rhBMP-2 treatment of MLB13MYC cells further supports the hypothesis that MLB13MYC are a useful model system with which to study the effects of rhBMP-2 and other growth and differentiation.

IT Animal cell line
(MLB13MYC, bone morphogenetic protein-2 effect on, cell
differentiation in relation to)

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